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**APTAMERS SELECTED FROM LIVE TUMOR CELLS  
AND THE USE THEREOF**

5 The present invention relates to aptamers selected from  
live tumor cells and to uses thereof in the diagnosis  
and treatment of certain cancers and other pathologies.

10 In the oncology field, non-invasive diagnostic methods  
by *in vivo* imaging (radiography, X-ray scanner, MRI,  
gamma-scintigraphy, positron emission tomography), are  
rarely specific for a molecular determinant (or marker)  
characteristic of the tumor to be diagnosed or to be  
treated. This contrasts with the precision of the  
knowledge obtained *in vitro*, which describes  
15 increasingly finely the molecular anomalies which cause  
cancerous processes, and results in the diagnostic  
tools being inappropriate for the current data of  
molecular science. The same dichotomy between *in vitro*  
and *in vivo* is often found in the field of anticancer  
20 therapies, which results in difficulties in developing  
treatments with an acceptable therapeutic index  
(effective dose/toxic dose).

The search for ligands capable of recognizing a  
25 molecular determinant (or marker) signaling a specific  
type of tumor, or else a given stage in its  
development, or alternatively signaling the metabolic  
state of a tumor, is therefore essential for better  
follow-up and better therapy of cancers. Unfortunately,  
30 these ligands are generally obtained from targets which  
have been purified and isolated out of their biological  
context, and which are therefore different from the  
targets placed in their natural environment.

35 Thus, most commonly, the ligands, which are effective  
in a test tube, are incapable of interacting with their  
target:

- because they cannot cross tissue barriers,
- because they are instable in the organism, or

responsible for too many adverse interactions with other biomolecules,

- because the natural structure of the target, placed out of its natural cell environment, was not conserved or because certain essential modifications of this structure, such as: (i) post-translational modifications of proteins or (ii) interactions with other proteins, cannot be reproduced *in vitro*.

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The latter two limitations are particularly frequent in the case of targets such as transmembrane proteins comprising a lipophilic segment inserted into the lipid cell membrane; this lipophilic segment is not conserved *in vitro*, whereas the membrane insertion of these proteins determines their structure and is essential to their activity.

Furthermore, the problem arises of the specificity with which the available ligands recognize the targets identified in tumors. It is therefore important to be able to provide specific ligands for diagnosing and/or treating certain cancers, in particular those related to the presence of a mutated tyrosine kinase receptor, resulting in constitutive activation or in over-expression of this receptor.

Molecular medicine therefore needs new molecular recognition probes which are:

- 30 - specific,
- adjustable, and
- easy to produce at a reasonable cost.

Pharmacological research has set up novel strategies for discovering novel ligands effective against targets identified in tumors:

- combinatorial libraries of small molecules make it possible to increase the chances of finding a ligand against a specific protein (example: a

subclass of combinatorial libraries, false substrates which inhibit enzymes, such as those which inhibit MMPs (*matrix metalloproteases*)). Their major drawback is that they are sorted *in vitro*. Moreover, their selectivity is not guaranteed, hence the difficulty in obtaining, with these agents, a compound which is sufficiently selective to be effective and free of side effects. Only exceptionally are compounds obtained which are specific for an abnormal protein form, which leads to non-specific binding and adverse effects and results in a poor therapeutic index;

- monoclonal antibodies are excellent agents for specific targeting and have recently been used for therapeutic purposes (example: Herceptin in breast cancer); however, they remain very difficult to use *in vivo* due to their sizes, *idiotypy* and immunogenicity, and are extremely expensive to produce and to optimize. Furthermore, monoclonal antibodies reach their limits when the recognition of a point mutation affecting a single amino acid on a protein is involved. In particular, no monoclonal antibody exists which is capable of identifying one of the abnormal forms of the Ret protein;

- aptamers, which constitute an alternative means of diagnosis and therapy, and which have a certain number of advantages compared with antibodies, as illustrated in Table I below.

	Aptamers	Antibodies
Size	9-15 kDa	>150 kDa
discrimination*	+++	+ / ++
affinity	5-100 nM	0.1-100 nM
Source	synthetic	animal
Cost	↘	↗

\*theophylline versus caffeine, for example.

A method of selecting aptamers, which bind specifically and have a high affinity for predefined targets, was described at the beginning of the 1990s, and known as the SELEX method (*Systematic Evolution of Ligands by Exponential enrichment*). This method operates in iterative cycles of selection-amplification; this method and a certain number of improvements and applications of this method are described in particular in the following American patents: US 5,270,163; 5 US 5,475,096; US 5,496,938; US 5,567,588; US 5,580,737; 10 US 5,637,459; US 5,660,985; US 5,683,867; US 5,707,796; US 5,763,177 and US 5,789,157.

Briefly, the principle of the SELEX method involves the 15 selection, from a mixture of nucleic acids comprising random sequences, and by successive reiterations of binding, separation and amplification steps, of nucleic acid molecules (aptamers) exhibiting a defined binding affinity and a defined specificity for a given target. 20 Thus, starting from a mixture of random nucleic acids, the SELEX method comprises more specifically the following steps:

- bringing the mixture of nucleic acids into contact with the target element (natural or synthetic 25 polymers: proteins, polysaccharides, glycoproteins, hormones, receptors, antigens, antibodies, cell surfaces; small molecules: medicament, metabolites, cofactors, substrates, transition state analogs; tissues and in 30 particular whole cells; viruses, etc.), under conditions which promote binding,
- separating the unbound sequences,
- dissociating the nucleic acid-target element complexes,
- 35 - amplifying the dissociated nucleic acids, so as to obtain a mixture enriched in ligands (aptamers), and
- repeating the binding, separation, dissociation and amplification steps, for the desired number of

times.

Among the abovementioned American patents:

- 5     -     US     patent     5,270,163     describes     the     initial  
principle of the SELEX method;
- 10    -     US     patent     5,580,737     describes     an     application     of  
this method, comprising a counterselection step,  
to the selection of aptamers capable of  
discriminating between two substances of similar  
15    structure, namely caffeine and theophylline;
- 15    -     US     patent     5,660,985     describes     an     application     of  
this method to the selection of modified aptamers  
containing modified nucleotides at the level of  
the pyrimidines (modification in the 5-position or  
15    in the 2'-position);
- 20    -     US     patent     5,789,157     describes     an     application     of  
this method to the selection of aptamers capable  
of binding to proteins present at the surface of  
tissues and in particular of cells; this patent  
20    recommends in particular applying the SELEX method  
to specific targets, namely tissues and in  
particular tumor cells, which are considered to be  
targets more complex than those used in the other  
patents cited, and which are generally targets  
25    that have been molecularly identified (proteins,  
etc.). This US patent 5,789,157 also specifies  
that it is possible to use a counter selection (or  
negative selection), before, during or after the  
implementation of the SELEX method. The negative  
30    selection more specifically described in US patent  
5,580,737 makes it possible to discriminate  
between tissue types which are different but  
nevertheless very similar. For example, negative  
selection can be used to identify ligands which  
35    exhibit a high specificity for tumor cells and not  
with respect to the corresponding normal cells.  
This patent also envisions the case where the  
nucleic acids which are ligands of a cell type  
which expresses a certain receptor can be

counterscreened with a cell line constructed in such a way that it does not express said receptor; however, the use of such a strategy can be difficult to implement if the protein induces a substantial phenotypic change in the cell;

- US patent 6,232,071 describes an application of this method to the selection of aptamers specific for tenascin C.

Surprisingly, the Applicant has found that it is possible to obtain aptamers specific for cell receptors, and more specifically for tumor markers, by carrying out the method known as SELEX on live target cells, under certain conditions.

Consequently, according to a first aspect, a subject of the invention is a method for identifying ligands or aptamers specific for a membrane receptor with tyrosine kinase activity (RPTK for *receptor protein-tyrosine kinase*) expressed in an activated form by cells (whatever the origin or the cause of the activation) or nonactivated form (preferably in an activated form), using a mixture of nucleic acids, which method comprises at least the following steps:

- (a) bringing a mixture of nucleic acids into contact with cells not expressing said receptor protein-tyrosine kinase or expressing it in a nonactivated form ( $C_N$  cells), said cells having the same cell type as cells expressing the same receptor protein-tyrosine kinase but in an activated form, due to the existence of at least one mutation in the extracellular domain ( $C_{Te}$  cells);
- (b) recovering a first subset  $S1$  of nucleic acids which do not bind to the  $C_N$  cells, in step (a);
- (c) bringing said first subset  $S1$  into contact with  $C_i$  cells, having the same cell type as the  $C_{Te}$  cells, but expressing said receptor protein-tyrosine kinase mutated in its intracellular part, said  $C_i$  cells exhibiting a phenotype of the same type as

- that of the  $C_{Te}$  cells;
- (d) recovering a second subset S2 of nucleic acids which do not bind to the  $C_i$  cells in step (c);
  - (e) bringing the second subset S2 into contact with the  $C_{Te}$  cells;
  - (f) recovering the nucleic acids which bind to said  $C_{Te}$  cells, i.e. those exhibiting a high affinity with respect to the cells expressing said receptor protein-tyrosine kinase mutated in the extracellular domain (activated receptor), after dissociation of the cell-nucleic acid complexes;
  - (g) amplifying said nucleic acids with high affinity for the cells expressing said receptor protein-tyrosine kinase mutated in the extracellular domain (activated receptor), so as to obtain a mixture of nucleic acids, enriched in nucleic acids having a high affinity for said  $C_{Te}$  cells, and
  - (h) identifying the ligands or aptamers specific for the cells expressing receptor protein-tyrosine kinases (RPTKs) in an activated form, from the mixture obtained in (g).

Surprisingly, such a method, even though it comprises steps for excluding aptamers which bind to nonactivated forms of RPTK, makes it possible to select aptamers specific for RPTK, i.e., either aptamers capable of binding to said RPTK and of inhibiting the activity of said RPTK (activation of the kinase cascade), or aptamers capable only of binding to said RPTK (of advantage in imaging applications).

#### Definitions

- Receptor protein-tyrosine kinases (RPTKs)

The receptor protein-tyrosine kinases (RPTKs) constitute a very large family of proteins. There are currently more than 90 known genes encoding protein

tyrosine kinases (PTKs), in the human genome (Blume-Jensen P. et al., Nature, 2001, 411, 355-365): 58 encode transmembrane RPTKs divided up into 20 families and 32 encode cytoplasmic PKTs. Among the human  
5 receptor protein-tyrosine kinases (RPTKs) involved in cancers, mention may be made of the following families: EGFR (Epithelial Growth Factor Receptor), InsulinR (Insulin Receptor), PDGFR (Platelet-derived Growth Factor Receptor), VEGFR (Vascular Endothelial Growth  
10 Factor Receptor), FGFR (Fibroblast Growth Factor Receptor), NGFR (Nerve Growth Factor Receptor), HGFR (Hepatocyte Growth Factor Receptor), EphR (Ephrin Receptor), AXL (Tyro 3 PTK), TIE (Tyrosine Kinase Receptor in endothelial cells), RET (Rearranged During  
15 Transfection), ROS (RPTK expressed in certain epithelial cells) and LTK (Leukocyte Tyrosine Kinase). In the subsequent text, the term "RPTK" used without other specification implies any receptor (activated or nonactivated; in an activated or nonactivated form).

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- Nonactivated RPTK

In its normal form, said receptor is not activated; activation is observed only after suitable stimulation  
25 of cells expressing said normal receptor (RPTK activated after stimulation).

30

- RPTK mutated in the extracellular domain (activated receptor)

In certain abnormal forms, due to mutations in the extracellular domain of the receptor protein-tyrosine kinase (one or more point mutations, insertions, deletions and/or rearrangements), a constitutive  
35 activation or an overexpression of the receptor is observed; such an activated receptor mutated in the extracellular portion is a constitutive activator of the kinase cascade.



- RPTK mutated in the intracellular domain

Such a receptor can activate certain intracellular cascades; it is not considered, for the purpose of the present invention, to be an activated receptor; on the other hand, it is included in the definition of receptors in or as an activated form.

- RPTK in or as an activated form

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The expression "receptor in or as an activated form" is intended to mean an RPTK which activates the kinase cascade, whatever the reason for this:

. activation by stimulation with a growth factor  
15 (normal activation, under certain conditions),  
. constitutive activation or overexpression of the receptor, due to the existence of one or more mutations, either in the extracellular portion, or in the intracellular portion of said receptor.

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Surprisingly, the specific conditions of the method according to the invention effectively make it possible to select and identify ligands or aptamers specific for the receptor protein-tyrosine kinase(s) preselected, i.e. which bind to said receptor, and in addition, among these, to select those capable of inhibiting said receptors in their activated form. In fact, the selection of the aptamers with  $C_N$ ,  $C_{Te}$  and  $C_i$  cells, as defined above, effectively makes it possible to obtain, in particular after repetition of steps a) to g), aptamers specific for cells expressing preselected receptor protein-tyrosine kinases (RPTKs).

30

In accordance with the invention, several cycles of steps (a) to (g) can advantageously be repeated using the mixtures enriched in ligands or aptamers from the preceding cycle, until at least one aptamer is obtained, the affinity of which, defined by its dissociation constant ( $K_d$ ), can be measured and is

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suitable for pharmaceutical use.

Also in accordance with the invention:

5 . the starting combinatorial library of nucleic acids advantageously consists of oligonucleotides comprising random sequences, of the same type as those described in the abovementioned SELEX patents. It contains at least  $10^2$  nucleic acids, preferably between  $10^9$  and  $10^{15}$  nucleic acids; the nucleic acids constituting said  
10 combinatorial library are preferably natural nucleic acid sequences (RNA or DNA) or modified nucleic acid sequences (for example, pyrimidines modified with a fluorine atom in the 2'-position of the ribose) consisting of random sequences comprising, respectively  
15 at their 5' and 3' ends, fixed sequences for PCR amplification, preferably the oligonucleotides of sequences SEQ ID NO: 1 and SEQ ID NO : 2, or a fragment of at least 8 nucleotides of these oligonucleotides. Said random sequences each contain between 10 and 1000  
20 nucleotides, preferably 50 nucleotides.

. the aptamers selected are defined by virtue of their primary sequence and by virtue of their secondary structure; the latter is either in hairpin loop form, or in the form of more complex structures (for example:  
25 pseudo-knot, triple helix, Guanine quartet, etc.). They advantageously consist preferably of nucleic acid sequences of 20 to 100 nucleotides.

Advantageously, the conditions described above make it  
30 possible to identify ligands or aptamers against molecular determinants (or markers) of pathologies, which can actually be effective under the very conditions of their future use, i.e. *in vivo*.

35 Thus, by selecting against a specific target, i.e. the  $C_N$ ,  $C_i$  and  $C_{Te}$  cells as defined above, aptamers are effectively obtained which specifically recognize cells expressing a preselected receptor protein-tyrosine kinase in particular in its activated form.

According to an advantageous embodiment of said method, the identification of the ligands or aptamers specific for the  $C_{Te}$  cells according to step (h) comprises an evaluation of the biological activity of said aptamers on said  $C_{Te}$  cells.

The biological activities, which are advantageously evaluated, depend on the receptor selected; they are in particular the following:

- (a) inhibition or activation of the autophosphorylation of the receptor (RPTK),
- (b) inhibition or activation of the kinase activation cascade,
- (c) inhibition of the phosphorylation of the normal RPTK of normal cells ( $C_N$  cells) activated by suitable stimulation (appropriate growth factor, for example),
- (d) reversion of the phenotype associated with activation of the RPTK.

For the purpose of the present invention, the following terms are considered to be equivalent: nucleic acid fragment, oligonucleotide, ligand or aptamer.

According to a second aspect, a subject of the present invention is ligands or aptamers, characterized in that they are specific for cells expressing a receptor protein-tyrosine kinase (RPTK) in an activated or nonactivated form (preferably in an activated form), in particular an RPTK mutated in the extracellular domain, and can be identified by means of the method for identifying aptamers, as defined above.

According to an advantageous embodiment of said aptamer, it is specific for cells expressing a receptor protein-tyrosine kinase (RPTK) in an activated or nonactivated form, selected in particular from the group consisting of the following membrane receptors,

given by way of nonlimiting examples: EGFR (Epithelial Growth Factor Receptor), InsulinR (Insulin Receptor), PDGFR (Platelet-derived Growth Factor Receptor), VEGFR (Vascular Endothelial Growth Factor Receptor), FGFR (Fibroblast Growth Factor Receptor), NGFR (Nerve Growth Factor Receptor), HGFR (Hepatocyte Growth Factor Receptor), EPHR (Ephrin Receptor), AXL (Tyro 3 PTK), TIE (Tyrosine Kinase Receptor in endothelial cells), RET (Rearranged During Transfection), ROS (RPTK expressed in certain epithelial cells) and LTK (Leukocyte Tyrosine Kinase).

According to an advantageous arrangement of this embodiment, said aptamer recognises in particular the Ret receptor activated by mutation at a cysteine located in the extracellular domain, preferably at codons 609, 611, 618, 620 or 634.

According to a preferred mode of this arrangement, said aptamer can be identified by means of a method, as defined above, which comprises:

- (a) bringing a mixture of nucleic acids into contact with  $C_N$  cells not expressing any Ret receptor in an activated form,
- (b) recovering a first subset  $S1$  of nucleic acids which do not bind to said  $C_N$  cells, in step (a),
- (c) bringing said first subset  $S1$  into contact with  $C_i$  cells expressing a Ret receptor, mutated in its intracellular domain, in particular the mutated receptor Ret<sup>M918T</sup>,
- (d) recovering a second subset  $S2$  of nucleic acids which do not bind to said  $C_i$  cells,
- (e) bringing the second subset  $S2$  into contact with  $C_{Te}$  cells expressing a Ret receptor activated by mutation in the extracellular domain, which receptor is selected from the group consisting of mutated Ret receptors carrying a mutation on one of the cysteines located in the extracellular domain, preferably at Cys609, Cys611, Cys618,

- Cys620 or Cys634, preferably the Ret<sup>C634Y</sup> receptor,
- (f) recovering the nucleic acids bound to said C<sub>Te</sub> cells, i.e. exhibiting both a high affinity and a binding specificity for the cells expressing a mutated Ret receptor (activated receptor) as defined in step (e),
- (g) amplifying said nucleic acids obtained in step (f), so as to obtain a mixture of nucleic acids, enriched in nucleic acids having a high affinity for the C<sub>Te</sub> cells,
- (h) repeating steps (a)-(g), until at least one aptamer is obtained, the affinity of which for the C<sub>Te</sub> cells, defined by its dissociation constant (K<sub>d</sub>), is measurable and suitable for a pharmacological activity, and
- (i) identifying the aptamers specific for the cells expressing a Ret receptor in its activated form, selected from the mixture obtained in (h).
- The cycle for obtaining the aptamers, according to the invention, applied to the Ret receptor, is illustrated in figure 1.

The Ret (rearranged during transfection) oncogene encodes an abnormal form of a receptor-type surface protein of the tyrosine kinase family; this protooncogene is located on chromosome 10q11.2. Mutations in the Ret protooncogene are associated with disparate diseases, in particular Hirschsprung's disease and multiple endocrine neoplasia type II (or MEN 2), which includes MEN type 2A (MEN 2A), MEN type 2B (MEN 2B) and familial medullary thyroid cancer (or FMTC). MEN 2A is characterized by a medullary thyroid carcinoma, a pheochromocytoma and parathyroid hyperplasia (primary hyperparathyroidism). MEN 2B is characterized by a particularly aggressive form of medullary thyroid cancer, a pheochromocytoma, multiple mucosal neurogliomas and intestinal ganglioneuromatosis. A truncated form of ret encodes an

intracellular protein associated with papillary thyroid cancer (PTC).

Oncogenes are mutated forms of protooncogenes, which  
5 are normal proteins, the function of which is to  
control cell growth and division, in particular after  
activation by appropriate growth factors (such as, for  
example, GDNF for the protooncogene encoding the Ret  
receptor). Certain mutations of these protooncogenes  
10 result in forms of these proteins which are permanently  
active, even in the absence of stimulation by the usual  
growth factor(s) (deregulation). This constitutive  
(permanent) activation results in a permanent  
stimulation of cell growth and division, and, in the  
15 end, in cancerization. The mutated form of the proto-  
oncogene is then referred to as a tumor-activating  
oncogene. Oncogenes can induce a cancerization, for  
example, by overproduction of growth factors, or by  
inundation of the cell with replication signals, or by  
20 uncontrolled stimulation of intermediate pathways, or  
by disorganized cell growth linked to a high level of  
transcription factors. Certain oncogenes are  
transmitted from generation to generation, when the  
protooncogene mutates in the germinal cells. This  
25 implies an inherited and dominant tumor predisposition.  
For example, multiple endocrine neoplasia type II (or  
MEN 2) is the result of a germinal transmission of the  
activated Ret oncogene.

30 Mutations in exons 10 and 11 of the protooncogene are  
observed in more than 95% of cases of MEN 2A and in  
more than 80% of cases of MTC; most of these mutations  
are located at five conserved cysteines located in the  
extracellular domain (codons 609, 611, 618, 620 and  
35 634). These mutants of the Ret receptor spontaneously  
form active homodimers at the surface of the cell,  
which induce morphological and biochemical changes,  
resulting in a pheochromocytoma-type phenotype,  
dependent on the Ret receptor in MEN2 syndromes;

mutations at codon 634 are the most frequent in MEN 2A. The activation of the Ret receptor can be followed by means of the phosphorylation cascade (Jhiang SM, Oncogene, 2000, 19, 5590-5597; Califano D et al., PNAS, 5 1996, 93, 7933-7937).

As regards MEN 2B, a point mutation in exon 16 of the protooncogene at codon 918 of the *ret* gene has been identified in approximately 95% of cases; this mutation 10 leads to the substitution of a threonine with a methionine, in the catalytic domain of the Ret receptor. This mutation results in activation of the receptor in the form of a monomer.

15 Surprisingly, the selection of the aptamers with  $C_N$ ,  $C_{Te}$  and  $C_i$  cells, as defined above, effectively makes it possible to obtain, after repetition of steps a) to g), aptamers specific for the human form of Ret receptors, and in particular Ret receptors in their activated 20 form, for example the mutated  $Ret^{C634Y}$  receptors, expressed by the  $C_{Te}$  cells, normal receptors, which may or may not be activated, or receptors mutated in the intracellular portion (see definitions).

25 In accordance with the invention:

- the  $C_N$  cells are in particular wild-type PC12 cells (reference ECACC No. 88022) or wild-type NIH 3T3 cells (reference ECACC No. 93061524),
- the  $C_i$  and  $C_{Te}$  cells are obtained by introducing an 30 oncogene bearing a mutation, respectively intracellular or extracellular (figure 2), in  $C_N$  cells in culture in such a way that the latter express the oncogene; in the context of the invention, a similar transformed phenotype (figure 35 3) is obtained in the two cases. The PC12 cells change from a relatively non-adherent round shape to an elongated shape with pseudoneurites which exhibit good adhesion capacities (Califano D et al., PNAS, 1996, 93, 7933-7937).

It is then possible to carry out the selection and counterselection on virtually identical phenotypes which differ only by virtue of the cellular location of the oncogenic mutation (figure 1); more specifically,  
5 the  $C_i$  cells obtained are called PC12/MEN 2B (or NIH/MEN 2B) cells and the  $C_{Te}$  cells obtained are called PC12/MEN 2A (or NIH/MEN 2A) cells.

10 Advantageously, the conditions described above make it possible to identify ligands or aptamers against molecular determinants (or markers) of pathologies, which will actually be effective under the very conditions of their future use, i.e. *in vivo*.

15 Thus, by selecting a specific target, i.e. the  $C_N$ ,  $C_i$  and  $C_{Te}$  cells, as defined above, aptamers are effectively obtained which specifically recognize the cells expressing the human form of the Ret receptor in  
20 an activated or nonactivated form, preferably in an activated form.

The identification of an aptamer specific for cells expressing a human form of the Ret receptor in its  
25 activated or nonactivated form, as defined above, advantageously comprises an additional step (j) consisting in evaluating its biological activity on said  $C_{Te}$  cells.

30 The biological activities which are advantageously evaluated are as follows:

- (a) inhibition or activation of the autophosphorylation of the Ret receptor,
- (b) inhibition or activation of the Erk kinase  
35 activation cascade, Erk being an intracellular protein downstream of Ret in the cascade,
- (c) inhibition of the phosphorylation of the normal Ret receptor of PC12 cells activated with GDNF and reversion of the phenotype which is associated



therewith,

(d) reversion of the phenotype associated with the activation of the RPTK overexpressing the oncogene (Ret<sup>C634Y</sup>).

5 Said aptamer can be obtained by means of a method of identification as specified above, and is selected from the group consisting of the aptamers of formula (I):



10

in which:

R<sub>1</sub> represents 5' GGGAGACAAGAAUAAACGCUCAA 3' (SEQ ID NO:1) or a fragment of 1 to 23 nucleotides of said SEQ ID NO:1;

15 R<sub>2</sub> represents 5' AACGACAGGAGGCUCACAACAGGA 3' (SEQ ID NO:2) or a fragment of 1 to 24 nucleotides of said SEQ ID NO:2, and

R represents a random sequence of 10 to 1000 nucleotides, preferably of 50 nucleotides.

20

According to an advantageous arrangement of this embodiment, R is preferably selected from the following sequences:

D4	5'GCGCGGGAUAGUAUGGAAGGAUACGUUAUACCGUGCAAUCCAGGGCAACG 3' (SEQ ID NO:3)
D12	5'GGGCUUCAUAAAGCUACACCGGCAACGCAGAAUAGCCUUAAGCCCGAGUU 3' (SEQ ID NO:4)
D14	5'GGCCAUAGCGCACCAACCAAGAGCAAAUCCCUAAGCGCGACUCGAGUGAGC 3' (SEQ ID NO:5)
D20	5'GGGCCAAUCGAAGCCGGUAAUUCCAAACUAAACGUGCAAACUGCACCCGC 3' (SEQ ID NO:6)
D24	5'GCGGUAUGUAGGGAUAGCACUUUUUUUGCGUAUACCUACACCGCAGCG 3' (SEQ ID NO:7)
D30	5'AGGCGAGCCCGACACGUCAGUAUGCUAGACAACAACGCCCGCGUGGUAC 3' (SEQ ID NO:8)
D32	5'CCCCGCUUUUUGACGUGAUCGAACGCGUAUCAGUAACGUCAGCAGUCGAGC 3' (SEQ ID NO:9)
D33	5'CAAAGCGUGUAUUCUGUGAGCCGACCAUCGUUGCGAACAUCCCCGGAACG 3' (SEQ ID NO:10)
D42	5'GACCCGUUAUGAAGGUGGCGCAGGACACGACCGUCUGCAAUUGAGCGAGC 3' (SEQ ID NO:11)
D60	5'CCGACCUGUACAGCAGUUAAGUUACACGUUUGAAACAACCGCGUUCGAGC 3' (SEQ ID NO:12)
D76	5'GGCUUACACGGAGAAACAAGAGAGCGGCCCAAACUUGAUUGACAGUGGCC 3' (SEQ ID NO:13)
D71	5'GGCCCUUAAACGCAAAAACGAAGGAUCAUCGAUUGAUCGCCUUAUGGGCU 3' (SEQ ID NO:14)
D87	5'CCGCGGUCUGUGGGACCCUUCAGGAUGAAGCGGCAACCAUGCGGGCC 3' (SEQ ID NO:15)

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The preferred aptamers in which the Rs are as defined above are represented by the sequences SEQ ID NO:22 (D4; figure 11), SEQ ID NO:25 (D24; figure 12), SEQ ID NO:31 (D30; figure 13), SEQ ID NO:32 (D12; figure 14),  
30 SEQ ID NO:33 (D71; figure 15).

In accordance with the invention, in said aptamers, the riboses of the purines bear, as is the case in natural RNA, a hydroxyl (OH) function on the carbon in the 2'-position, while the riboses of the pyrimidines bear a fluorine atom on the carbon in the 2'-position. This modification of the 2'-position is known to confer on the nucleic acids a greater resistance with respect to nucleases.

10 The sequences of the primers used to carry out step (g) consisting in amplifying the mixture of nucleic acids of formulae  $R_1$ -R-R $_2$ , in which  $R_1$  represents SEQ ID NO:1 and  $R_2$  represents SEQ ID NO:2, are advantageously as follows:

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- Sense primer (primer P10):  
TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA (SEQ ID NO:16);

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- Antisense primer (primer P30):  
TCCTGTTGTGAGCCTCCTGTCGTT (SEQ ID NO:17).

The prediction of secondary and tertiary structure of the aptamers selected is carried out using the  
25 **RNAstructure** software written by David H. Mathews:  
<http://rna.chem.rochester.edu>. The algorithm used by this software is based on the searches described in the publication: D.H. Mathews et al., J. Mol. Biol., 1999, 288, 911-940. The same predictions can be obtained  
30 using the **mfold** algorithm, available of the site of the Michael Zuker laboratory:  
<http://bioinfo.math.rpi.edu/~zukerm/>. The algorithm used by this software is also based on the searches described in the publication D.H. Mathews et al.,  
35 mentioned above.

Among the aptamers described above, some have a common structure, defined by formula II below:

5'R<sub>4</sub>X<sub>6</sub>X<sub>5</sub>X<sub>4</sub>X<sub>3</sub>GGAAUAGX<sub>2</sub>X<sub>1</sub>R<sub>3</sub>X'<sub>1</sub>X'<sub>2</sub>CGUAUACX'<sub>3</sub>X'<sub>4</sub>X'<sub>5</sub>X'<sub>6</sub>R<sub>5</sub> 3' (II),

the secondary structure of which is represented in figure 10, and in which:

- the riboses of the purines bear an OH group in the 2'-position and the riboses of the pyrimidines bear a fluorine atom in the 2'-position,
- **R<sub>3</sub>** is present or absent and represents an apical bulge (or loop) comprising:
  - . a linear or branched carbon chain selected from the group consisting of C<sub>6</sub>-C<sub>30</sub> alkyl groups or C<sub>6</sub>-C<sub>30</sub> aryl groups;
  - . a polymer such as PEG or PEI, or the like;
  - . functional groups such as biotin, streptavidin, peroxidase, etc.;
  - . other molecules of interest such as, for example, active ingredients, labeling tags, in particular fluorescent tags, or chelating agents for radioisotopes;
  - . a natural (DNA or RNA) or modified nucleotide sequence (for example: 2'-fluoro, 2'-O-methyl, PNA, LNA, etc.); preferably, **R<sub>3</sub>** represents the following bulges or loops (1) to (4):
    - loop (1): 5' UGGAAGGA 3' (SEQ ID NO:29)
    - loop (2): 5' CUUUUUU 3' (SEQ ID NO:30)
    - loop (3): 5' GNPuA 3'
    - loop (4): 5' UNCG 3',
- in which the riboses of the purines bear a hydroxyl function on the carbon in the 2'-position, while the riboses of the pyrimidines bear a fluorine atom on the carbon in the 2'-position,
- **X<sub>1</sub>, X'<sub>1</sub>, X<sub>2</sub>, X'<sub>2</sub>, X<sub>3</sub>, X'<sub>3</sub>, X<sub>4</sub>, X'<sub>4</sub>, X<sub>5</sub>, X'<sub>5</sub>, X<sub>6</sub> and X'<sub>6</sub>** represent Py or Pu with, preferably:
  - X<sub>1</sub>-X'<sub>1</sub> corresponding to C-G, A-U, G-C or U-A
  - X<sub>2</sub>-X'<sub>2</sub> corresponding to C-G, A-U, G-C or U-A
  - X<sub>3</sub>-X'<sub>3</sub> corresponding to C-G, A-U, G-C or U-A
  - X<sub>4</sub>-X'<sub>4</sub> corresponding to C-G, A-U, G-C or U-A
  - X<sub>5</sub>-X'<sub>5</sub> corresponding to C-G, A-U, G-C or U-A
  - X<sub>6</sub>-X'<sub>6</sub> corresponding to C-G, A-U, G-C or U-A
  - N corresponding to G or C or A or U,
  - Pu** corresponding to G or A, in which the riboses bear

an OH group in the 2'-position (natural RNA chemistry),  
**Py** corresponds to U or C, in which the riboses bear a  
fluorine atom in the 2'-position, and

- **R<sub>4</sub>** and **R<sub>5</sub>** are present or absent and represent:

- 5 . a natural (DNA or RNA) or modified nucleotide  
sequence (for example: 2'-fluoro, 2'-O-methyl, PNA,  
LNA, etc.), comprising between 1 and several  
thousand nucleotides, preferably between 1 and 39  
nucleotides; a part of said nucleotide sequence or  
10 said sequence preferably comprising one of the  
following sequences:

**R<sub>4</sub>** :

5'-**R<sub>1</sub>**-**Z<sub>1</sub>**-3', with **Z<sub>1</sub>**=G: 5' GGGAGACAAGAAUAAACGCUCAAG 3'  
(SEQ ID NO:18) or

- 15 5'-**R<sub>1</sub>**-**Z<sub>1</sub>**-3', with **Z<sub>1</sub>**=GCGGUAAU (SEQ ID NO:26):

5' GGGAGACAAGAAUAAACGCUCAAGCGGUAAU (SEQ ID NO:19), and

**R<sub>5</sub>** :

5'-**Z<sub>2</sub>**-**R<sub>2</sub>**-3', with **Z<sub>2</sub>**=CAAUCCAGGGCAACG (SEQ ID NO:27):

- 20 5' CAAUCCAGGGCAACGAACGACAGGAGGCUCACAAACAGGA 3' (SEQ ID  
NO:20) or

5'-**Z<sub>2</sub>**-**R<sub>2</sub>**-3', with **Z<sub>2</sub>**=ACCGCAGCG (SEQ ID NO:28):

5' ACCGCAGCGAACGACAGGAGGCUCACAAACAGGA 3' (SEQ ID NO:21),

- 25 . a linear or branched carbon chain selected from the  
group consisting of C<sub>6</sub>-C<sub>30</sub> alkyl groups or C<sub>6</sub>-C<sub>30</sub> aryl  
groups;

- 30 . a polymer such as PEG or PEI, or the like,  
functional groups such as biotin, streptavidin,  
peroxidase, etc.,  
other molecules of interest such as, for example,  
active ingredients, labeling tags, in particular  
fluorescent tags, or chelating agents for  
radioisotopes.

35 Among the aptamers having a structure as defined in  
formula II and in figure 10, differences in properties  
as specified hereinafter are observed:

The structure of formula II, in which **R<sub>3</sub>**, **R<sub>4</sub>** and **R<sub>5</sub>** are  
absent, is sufficient for binding to the Ret<sup>C634Y</sup>

receptor, whereas the structure of formula II, in which R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> are present, exhibit either binding properties only, or both binding and inhibiting properties; these properties varying as a function of R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub>:

- When R<sub>3</sub> represents 5' UGGAAGGA 3' (SEQ ID NO:29: loop (1)), R<sub>4</sub> represents SEQ ID NO:18 and R<sub>5</sub> represents SEQ ID NO:20, the aptamer exhibiting such a structure (family D4) has both properties of binding to said Ret receptor in its activated form and properties of inhibition of the activity of said receptor. The secondary structure of this product is represented in figure 11; it involves the products of family D4, in which R<sub>4</sub> and R<sub>5</sub> comprise a sufficient number of nucleotides resulting in inhibition of the activity of the Ret receptor. A preferred aptamer corresponding to this definition comprises, from 5' to 3', successively the following sequences: SEQ ID NO:1+SEQ ID NO:3+SEQ ID NO:2 (SEQ ID NO:22) (D4: figure 5A and figure 11), with reference to formula I.

- When R<sub>3</sub> represents 5' CUUUUUU 3' (SEQ ID NO:30: loop (2)), 5'GNPuA 3' (loop 3) or 5'UNCG 3' (loop 4), R<sub>4</sub> comprises from 1 to 30 nucleotides selected from SEQ ID NO:19 or from 1 to 24 nucleotides selected from SEQ ID NO:18 and R<sub>5</sub> comprises from 1 to 33 nucleotides selected from SEQ ID NO:21 or from 1 to 39 nucleotides selected from SEQ ID NO:20, the aptamer exhibiting such a structure has only properties of binding to said Ret receptor in its activated form, and in particular to the Ret receptor mutated in its extracellular domain. When R<sub>3</sub> represents 5' CUUUUUU 3' (loop 2), R<sub>4</sub> represents SEQ ID NO:19 and R<sub>5</sub> represents SEQ ID NO:21, the secondary structure of this product is represented in figure 12; the preferred product is represented by SEQ ID NO:25 and belongs to the family D24, in which R<sub>4</sub> and R<sub>5</sub> comprise at least one nucleotide. This aptamer comprises successively from 5' to 3': SEQ ID NO:1 + SEQ

ID NO:7 + SEQ ID NO:2, with reference to formula I.

5     - When R<sub>3</sub> represents 5' UGGAAGGA 3' (loop 1), and in the absence of R<sub>4</sub> and of R<sub>5</sub>, the aptamer exhibiting such a structure has only properties of binding to said Ret receptor in its activated form. A preferred aptamer corresponding to the latter definition corresponds to a part of SEQ ID NO:3 of family D4 (SEQ ID NO:23).

10    The differences between the definitions of R<sub>1</sub> and R<sub>2</sub> in formula I R<sub>1</sub>-R-R<sub>2</sub> and of R<sub>4</sub> and R<sub>5</sub> in formula II come from the non-superposition between the consensus sequence of figure 10 and the definition of R (random sequences) in formula I.

15

Figures 5A and 10 to 12 show the links between the two formulae (I and II).

20    The ligands or aptamers according to the invention can advantageously be used in the following applications:

25       - as diagnostic reagent: both *in vitro* (diagnostic kits; histological marker; chips) and *in vivo* (contrast agent for imaging; radiopharmaceuticals). In fact, the aptamers according to the invention which have an ability to bind specifically to a Ret receptor in its activated or nonactivated form are suitable for the detection of cells expressing said receptor in activated or nonactivated form; in particular, when the Ret receptor is mutated in its extracellular domain,  
30    said aptamers are particularly suitable for the detection of abnormal cells expressing said receptor; when the receptor in its activated form is not mutated, a difference in level of expression of the protein will then be followed;

35       - as medicament, in particular as anticancer agent; in particular as regards the aptamers which exhibit both an ability to bind to the Ret receptor and an inhibitory action with respect, in particular, to the mutated Ret<sup>C634Y</sup> receptor.

In fact, surprisingly:

- some of the aptamers selected have only a binding activity and will preferably be used as diagnostic reagents;

- other aptamers selected also exhibit an inhibitory activity on the oncogenic transformation mediated by the human Ret oncogene mutated in particular at a cysteine in the extracellular domain, and an inhibitory effect on the activity of the normal Ret protein after stimulation by its own ligand (family D4 or sequences derived from formula II with chemical modifications). The aptamers exhibiting such an activity may be used either as diagnostic agents or as medicaments. In order to characterize them, four tests can advantageously be carried out, as specified below:

(a) inhibition or activation of the autophosphorylation of Ret,

(b) inhibition or activation of the Erk kinase activation cascade, Erk being an intracellular protein downstream of Ret in the cascade,

(c) inhibition of Ret phosphorylation activated by GDNF,

(d) reversion of the phenotype associated with the activation of Ret (in particular, Ret<sup>C634Y</sup>).

A subject of the present invention is also a reagent for diagnosing a tumor, characterized in that it consists of at least one aptamer as defined above.

According to an advantageous embodiment of said reagent, it corresponds to an aptamer of formula II, as defined above:

5' R<sub>4</sub> X<sub>6</sub> X<sub>5</sub> X<sub>4</sub> X<sub>3</sub> GGAUAGX<sub>2</sub> X<sub>1</sub> R<sub>3</sub> X'<sub>1</sub> X'<sub>2</sub> CGUAUACX'<sub>3</sub> X'<sub>4</sub> X'<sub>5</sub> X'<sub>6</sub> R<sub>5</sub> 3',

in which R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> are absent.

According to an advantageous arrangement of this

embodiment, said reagent corresponds to an aptamer of sequence:

5'GUAGGGAAUAGCACGUAUACCUAC3' (SEQ ID NO:24),

in which  $X_1-X'_1 = A-U$ ,  $X_2-X'_2 = C-G$ ,  $X_3-X'_3 = G-C$ ,  $X_4-X'_4 =$   
5  $A-U$ ,  $X_5-X'_5 = U-A$  and  $X_6-X'_6 = G-C$ .

According to another advantageous embodiment of said reagent, it corresponds to an aptamer of formula II, in which  $R_3$  represents 5' CUUUUUU 3' (loop (2)),  $R_4$   
10 represents the sequence SEQ ID NO:19 and  $R_5$  represents the sequence SEQ ID NO:21; this aptamer corresponds to SEQ ID NO:25, and comprises successively from 5' to 3', with reference to formula I: SEQ ID NO:1 + SEQ ID NO:7 + SEQ ID NO:2, as specified above.

15 A subject of the present invention is also a reagent for diagnosing or detecting the Ret receptor in an activated or nonactivated form, characterized in that it consists of at least one aptamer as defined above.

20 A subject of the present invention is also a medicament, characterized in that it comprises an aptamer as defined above which has both an ability to bind to an RPTK receptor and an inhibitory action with  
25 respect to said receptor in an activated form.

A subject of the present invention is also a medicament for use in the treatment of a tumor, characterized in that it comprises an aptamer as defined above, which  
30 has both an ability to bind to an activated RPTK receptor, and in particular to a receptor mutated in the extracellular domain, and in particular to the Ret receptor mutated, for example, at one of the cysteines located in the extracellular domain (codons 609, 611,  
35 618, 620 and 634), and an inhibitory action with respect to this activated receptor.

According to an advantageous embodiment of said medicament, it corresponds to an aptamer of the aptamer



family D4, as defined above.

5 A subject of the present invention is also a pharmaceutical composition, characterized in that it comprises an aptamer as defined above, which has both an ability to bind to an RPTK receptor and an inhibitory action with respect to said receptor in its activated form.

10 A subject of the present invention is also a pharmaceutical composition, characterized in that it comprises:

- 15 - an aptamer as defined above, which has both an ability to bind to an RPTK receptor mutated in the extracellular domain, and in particular to the Ret receptor mutated, for example, at one of the cysteines located in the extracellular domain (codons 609, 611, 618, 620 and 634), and an inhibitory action with respect to this mutated receptor,
- 20 - another anticancer molecule, and
- at least one pharmaceutically acceptable vehicle.

25 A subject of the present invention is also the use of an aptamer which has both an ability to bind to an RPTK receptor and possibly an inhibitory action with respect to this RPTK receptor, for screening products which interact with the RPTK receptor and which may or may not inhibit it.

30 A subject of the present invention is also the use of an aptamer which has both an ability to bind to an RPTK receptor in its activated form, and in particular to the Ret receptor mutated at one of the cysteines located in the extracellular domain (codons 609, 611, 35 618, 620 and 634), and possibly an inhibitory action with respect to this mutated RPTK receptor, for screening products which interact with the RPTK receptor and which may or may not inhibit it.

A subject of the present invention is also a method for screening products which interact with an RPTK receptor or targets which form a complex with the RPTK (in an activated or nonactivated form), which method is characterized in that it comprises:

- bringing cells expressing RPTKs in an activated or nonactivated form into contact with the substance to be tested,
- adding, under suitable conditions, an optionally labeled aptamer, as defined above, before the substance to be tested, at the same time or after it, and
- evaluating the competitive binding between the aptamer and the substance to be tested (for example: by measuring radioactivity, fluorescence, luminescence, surface plasmon resonance, BRET, FRET, or any other technique for demonstrating a molecular interaction).

In accordance with the invention, after identification of the substances which bind competitively with the aptamer to the cells exhibiting RPTKs in an activated form, the effect of these substances on the biological activity of said cells can be evaluated in order to find substances which inhibit or activate said biological activities of the cells expressing RPTKs in an activated form.

Besides the above arrangements, the invention also comprises other arrangements, which will emerge from the following description, which refers to examples of implementation of the method which is the subject of the present invention and also to the attached drawings, in which:

- Figure 1: diagrammatic selection of aptamers specific for PC12 MEN 2A cells:

. steps a) and b): counterselection

A combinatorial library of 2'-F-Py RNAs is incubated

with wild-type PC12 cells (PC12 wt) in suspension; the sequences not bound are recovered by centrifugation and incubated with PC12 MEN 2B cells; the sequences not bound, present in the supernatant, are recovered and  
5 incubated with PC12 MEN 2A cells.

. step c): selection

The unbound sequences are removed by means of several washes of the cells and the bound sequences are  
10 recovered by extraction with phenol.

. steps d) and e): amplification

The sequences selected are amplified by RT-PCR and in  
15 vitro transcription before a further selection cycle.

- Figure 2: structure of the various Ret receptors: normal Ret; MEN 2A Ret (Ret<sup>C634Y</sup>); MEN 2B Ret (Ret<sup>M918T</sup>).  
20

- Figure 3: representation of the phenotype of PC12 cells stably transfected with an expression vector containing the sequence encoding the human mutated receptor Ret<sup>C634Y</sup> (PC12 MEN 2A) or the mutated receptor  
25 Ret<sup>M918T</sup> (PC12/MEN 2B).

- Figure 4: diagrammatic representation of GDNF-dependent Ret receptor activation.

30 - Figure 5 (A): comparison of the prediction of the secondary structure of the D4 and D24 aptamers. The secondary structure prediction is carried out using the **RNAstructure** software written by David H. Mathews, <http://rna.chem.rochester.edu>. The algorithm is based  
35 on the searches described in D.H. Mathews et al. (Journal of Molecular Biology, 1999, 288, 911-940, mentioned above).

The same predictions can be obtained using the **mfold**

algorithm, available on the site  
<http://bioinfo.math.rpi.edu/~zukerm/>. The latter  
algorithm is based on the searches described in the  
publication in the name of D.H. Mathews et al.,  
5 mentioned above. The consensus structure is in bold  
characters. (B): curve of binding of the D4 aptamer  
with PC12 MEN 2A cells; the D4 aptamer is radiolabeled  
with  $^{32}\text{P}$  and incubated at various concentrations with  
cell monolayers. After several washes, the bound  
10 aptamer is quantified. The background noise is taken  
into account by subtracting, for each point obtained,  
the value obtained with a destructured D4 aptamer  
(D4Sc) having a scrambled sequence (i.e. containing the  
same nucleotides, but in a different order). A  
15 Scatchard analysis (insert) is used to evaluate the  
binding constant and the number of targets.

- Figure 6: effects of the various selected  
aptamers on the activity of the Ret<sup>C634Y</sup> receptor: (A)  
20 the PC12 MEN 2A cells are either nontreated, or treated  
for 16 hours with 150 nM of the aptamer indicated or of  
the starting RNA pool (combinatorial library); (B) the  
PC12 MEN 2A cells are treated for one hour with  
increasing doses of D4 (product of formula II) (left)  
25 or with 200 nM of product D4 for the incubation times  
indicated (right). The cell lysates are subjected to  
analysis by immunoblotting with anti-Ret (Tyr-  
phosphorylated) antibodies or anti-(phospho) Erk  
antibodies, as indicated. In order to confirm equal  
30 load, the blotting membranes are subjected to a further  
analysis in the presence of the abovementioned anti-  
total Ret and anti-total Erk antibodies. The nontreated  
control cells are indicated by a "C". The  
phosphorylation values, taking the value 1 for the  
35 control, were calculated using the NIH Image program,  
based on the sum of the two bands specific for Erk. The  
standard deviations are obtained from four independent  
experiments.

- Figure 7: effect of the D4 aptamer on the activity of the wild-type Ret receptor (Ret<sup>wt</sup>) and of the mutated Ret<sup>M918T</sup> receptor. (A) The PC12 cells transfected so as to stably express the nonmutated Ret receptor (PC12/wt) are treated for 10 min with GDNF (50 ng/ml) and soluble GFR $\alpha$ 1 (1.6 nM), or 5 min with NGF (100 ng/ml) and also with, simultaneously, 200 nM either of D4 aptamer or of the starting RNA pool. (B) The PC12 MEN 2B cells are serum-deprived for 6 hours and then treated for 1 hour with 200 nM of D4 aptamer or of starting RNA pool. The cell lysates are analyzed by immunoblotting with the following antibodies: anti-Ret (Tyr-phosphorylated) antibodies or anti-(phospho) Erk antibodies, as indicated. The standard deviations are obtained from five independent experiments.

- Figure 8: the D4 aptamer inhibits the differentiation of PC12 cells transfected so as to stably express the nonmutated Ret receptor and the GFR $\alpha$ 1 coreceptor (PC12- $\alpha$ 1/wt) induced by GDNF. The cells are either nonstimulated (A) or stimulated with GDNF alone (B) or stimulated with GDNF in the presence of the D4 aptamer or of the destructured D4 aptamer (D4Sc) (C and D, respectively). After treatment for 48 hours with GDNF, the percentage extension of the processes is calculated. The data are expressed as the percentage of cells comprising processes relative to the total number of cells counted. Each experiment is repeated at least three times (E) and the cell lysates are analyzed by immunoblotting with anti-VGF antibodies, VGF being a marker for differentiation induced by GDNF-induced Ret activation (F).

- Figure 9: the D4 aptamer modifies the morphology of the transformed NIH/MEN 2A cells. The cell lines indicated are seeded at equal density onto culture plates comprising 12 wells. One day after seeding, 3  $\mu$ M of D4 or of destructured D4 are added to the medium and the cells are maintained in culture for

72 hours, adding 3  $\mu$ M of each aptamer every 24 hours. Given the half-life of the aptamer in 15% serum, this protocol ensures the continuous presence of at least 200 nM of aptamer in the medium. The cells are  
5 photographed using a phase-contrast microscope.

- Figures 10 to 15: secondary structure of the following aptamers: formula II (figure 10); D4 (figure 11); D24 (figure 12); D30 (figure 13); D12 (figure 14)  
10 and D71 (figure 15).

- Figure 16: screening of aptamers which interact with RET on PC12 MEN 2A cells, by competitive binding with the D4 aptamer. The D4 aptamer is radiolabeled  
15 with  $^{32}$ P and incubated at 50 nM with monolayers of PC12 MEN 2A cells in the presence of 400 nM of various aptamers. After several washes, the amount of D4 aptamer bound is quantified. The background noise is taken into account by subtracting, for each point  
20 obtained, the value obtained with a destructured D4 aptamer (D4Sc) having a scrambled sequence (i.e. containing the same nucleotides, but in a different order).

- Figure 17: competitive binding of the E38 aptamer on PC12 MEN 2A cells in the presence of a range of D4 aptamer. The E38 aptamer is radiolabeled with  $^{32}$ P and incubated at 100 nM with monolayers of PC12 MEN 2A cells in the presence of an increasing concentration of  
30 D4 aptamer. After several washes, the amount of E38 aptamer bound is quantified. The background noise is taken into account by subtracting, for each point obtained, the value obtained with a destructured D4 aptamer (D4Sc) having a scrambled sequence (i.e.  
35 containing the same nucleotides, but in a different order).

It should be clearly understood, however, that these examples are given only by way of illustration of the

subject matter of the invention, of which they in no way constitute a limitation.

**EXAMPLE 1: Preparation of a combinatorial library of 2'-F-Py RNAs**

In order to obtain the 2'-F-Py RNA, it is necessary to carry out an *in vitro* transcription from a double-stranded DNA template obtained according to 3 methods:

**1. Before selection, by PCR amplification of the DNA sequence:**

B2S0: 5'TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCC3'

where N represents a random sequence of 50 nucleotides

**1st PCR cycle:**

*\* Hybridization*

5'TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCC3' (B2S0)

(primer P10)

3'AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'

(SEQ ID NO:16)

*\* Elongation*

5'TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCCTATAGTGAGTCGTATTA3'

3'**AGGACAACACTCGGAGGACAGCAA-M-AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'**,

where the text in bold represents the polymerized sequence and M represents the sequence complementary to N.

**2nd PCR cycle:**

*\* Denaturation*

5'TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCCTATAGTGAGTCGTATTA3'

3'**AGGACAACACTCGGAGGACAGCAA-M-AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'**

*\* Hybridization*

5'TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCCTATAGTGAGTCGTATTA3'

3'AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'

(primer P10: SEQ ID NO:16)

5'TCCTGTTGTGAGCCTCCTGTCGTT3' (primer P30 : SEQ ID NO :17)

3'AGGACAACACTCGGAGGACAGCAA-M-AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'

*\* Elongation*

5' TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCCTATAGTGAGTCGTATTA3'  
3' AGGACAACACTCGGAGGACAGCAA-M-AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'

5' TCCTGTTGTGAGCCTCCTGTCGTT-N-TTGAGCGTTTATTCTTGTCTCCCTATAGTGAGTCGTATTA3'  
3' AGGACAACACTCGGAGGACAGCAA-M-AACTCGCAAATAAGAACAGAGGGATATCACTCAGCATAAT5'

the text in bold representing the polymerized sequence and M represents the sequence complementary to N.

5

This second PCR cycle is repeated 15 to 30 times in order to obtain a double-stranded DNA which will be transcribed, *in vitro*, into 2'-F-Py RNA.

10 **2. During selection, by RT-PCR amplification of the selected 2'-F-Py RNAs, of formula R<sub>1</sub>-R-R<sub>2</sub>, as defined above:**

5' GGGAGACAAGAAUAAACGCUCAA-R-AACGACAGGAGGCUCACAACAGGA3',  
where R represents the sequence of the 2'-F-Py RNAs  
15 selected.

**Reverse transcription (RT):**

*\* Hybridization*

5' GGGAGACAAGAAUAAACGCUCAA-R-AACGACAGGAGGCUCACAACAGGA3' (2'-F-Py RNA)  
3' TTGCTGTCCTCCGAGTGTGTCCT5' (primer P30)

*\* Elongation*

20 5' GGGAGACAAGAAUAAACGCUCAA-R-AACGACAGGAGGCUCACAACAGGA3'  
3' **CCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTGTCCT5'**,

where the text in bold represents the polymerized sequence and S represents the sequence complementary to R.

25 **1st PCR cycle:**

*\* Denaturation*

3' **CCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTGTCCT5'**  
(cDNA of the 2'-F-Py RNA)

30

*\* Hybridization*



5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA3' (primer P10)  
3' CCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

\* *Elongation*

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

where the text in bold represents the polymerized sequence and S represents the sequence complementary to R.

5

**2nd PCR cycle:**

\* Denaturation

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'

3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

10

\* Hybridation

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
(primer P30) 3' TTGCTGTCCTCCGAGTGTTCCT5'

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA3' (primer P10)  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

\* Elongation

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

15 where the text in bold represents the polymerized sequence and S represents the sequence complementary to R.

20 This second PCR cycle is repeated 15 to 30 times in order to obtain a double-stranded DNA which will be transcribed, *in vitro*, into 2'-F-Py RNA.

**3. After selection, by PCR amplification of the aptamers from plasmid in which they have been cloned**

25 The plasmids contain the sequence:

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

where R represents the DNA sequence specific for the aptamer and S the sequence complementary to R.

5

**1st PCR cycle:**

*\* Denaturation*

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

10

*\* Hybridization*

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAAACAGGA3'  
(primer P30) 3' TTGCTGTCCTCCGAGTGTTCCT5'

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA3' (primer P10)  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

15

*\* Elongation*

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTCCT5'

where the text in bold represents the polymerized sequence and S represents the sequence complementary to

20

R.

This PCR cycle is repeated 15 to 30 times in order to obtain a double-stranded DNA which will be transcribed, *in vitro*, into 2'-F-Py RNA.

25

**in vitro transcription:**

One of the two strands of the PCR-amplified DNA serves as a template for the *in vitro* transcription of the double-stranded 2'-F-Py RNAs. The sequence underlined corresponds to the region of the T7 phage RNA polymerase promoter.

30

5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTGTCCT5'  
/ GGGAGACAAGAATAAACGCTCAA-R-AACGACAGGAGGCTCACAACAGGA3'  
5' TAATACGACTCACTATAGGGAGACAAGAAUAAACGCUCAA-R-AACGACAGGAGGCUACACAACAGGA3'  
3' ATTATGCTGAGTGATATCCCTCTGTTCTTATTTGCGAGTT-S-TTGCTGTCCTCCGAGTGTTGTCCT5'

In the 2'-F-Py RNAs, the sequence complementary to the  
5 primer P30 is at the 3' end:

5'AACGACAGGAGGCUACACAACAGGA3' (R<sub>2</sub>=SEQ ID NO:2)

and a part of sequence identical to the primer P10 is  
at the 5' end:

5'GGGAGACAAGAAUAAACGCUCAA3' (R<sub>1</sub>= SEQ ID NO:1).

10

## **EXAMPLE 2 : Materials and methods**

### *- Cell culture and immunoblotting analysis*

The conditions for growth of the PC12 cells and of the  
derived cell lines were described by D'Alessio A. et  
15 al. (Endocrinology 2003; 144, 10, 4298-4305).

The NIH/MEN2A and NIH/MEN2B cells are obtained from NIH  
3T3 cells stably transfected with expression vectors  
for Ret<sup>C634Y</sup> and Ret<sup>M918T</sup>. In order to evaluate the  
20 effects of the aptamers according to the invention on  
the Ret activity, cells (160 000 cells/3.5 cm of plate)  
were serum-deprived for 2 hours, and then treated with  
the amount indicated in the figures (200 nM) of aptamer  
or of a pool of primary RNA after a short denaturation-  
25 renaturation step.

When it is indicated, 100 ng/ml of 2.5 S NGF (Nerve  
Growth Factor, Upstate Biotechnology Inc., Lake  
Placid), 50 ng/ml of GDNF (Promega) or 1.6 nM of GFR $\alpha$ 1-  
30 FC chimera (R&D Systems Ltd., UK) are added to the  
culture medium.

The cell extracts and the immunoblotting analysis are  
carried out as described in Cerchia L. et al. (Biochem.  
35 J. 2003, 372, 897-903).

The primary antibodies used are as follows: anti-Ret antibody (C-19), anti-VGF antibody (R-15), anti-ERK1 antibody (C-16) (Santa Cruz Biotechnology Inc., Santa Cruz CA), anti-Ret (Tyr phosphorylated) antibody (Cell Signaling), anti-phospho44/42 MAP kinase monoclonal antibodies (E10) (Cell Signaling). For the immunoblots illustrated in the figures, the statistical analysis was carried out on at least four independent experiments.

- Cell process extension assay

PC12- $\alpha$ 1/wt cells are seeded at equal density onto culture plates comprising 12 wells. In order to evaluate the effects of the D4 aptamer on cell differentiation, the cells are pretreated for 6 hours with 400 nM of D4 aptamer or of destructured D4 aptamer, and then incubated with 50 ng/ml of GDNF and the appropriate aptamers at a final concentration of 3  $\mu$ M. After stimulation with GDNF for 24 hours, 3  $\mu$ M of D4 aptamer or of destructured D4 aptamer are again added to the cells and the stimulation is pursued up to 48 hours. At least 15 random fields are photographed 24 hours and 48 hours after the stimulation with GDNF, using a phase-contrast microscope, and 50 cells per frame are counted; the presence or the absence of process extension is recorded. It is considered that process extension exists when an extension process having a diameter more than double the diameter of the cell body is observed.

- SELEX ex vivo

The SELEX cycle is carried out essentially as described previously (Tuerk C et al., Science 1990, 249, 4968, 505-510; Ellington AD et al., Nature, 1990, 346, 6287, 818-22). The transcription is carried out in the presence of 1 mM of 2'-F-pyrimidines and of a mutant form of T7 RNA polymerase (T7<sup>Y639F</sup>) (Padilla, R et al., Nucleic Acids Res, 1999, 27, 6, 1561-1563), in order to

increase the yields. The 2'-F-Py RNAs are used because of their resistance to degradation by serum nucleases.

The complexity of the initial sample is approximately  
5  $10^{14}$  different sequences. The 2'-F-Py RNA library (1-5 nmol) containing 50 nucleotides of random sequences is heated at 85°C for 5 min in 3 ml of RPMI 1640, rapidly cooled in ice, for 2 min, and then reheated up to 37°C, before incubation thereof with the cells.

10

Two counterselection steps are carried out at each cycle.

In order to avoid the selection of aptamers which  
15 recognize the cell surface non-specifically, the combinatorial library of initial RNAs is first incubated for 30 minutes at 37°C with  $5 \times 10^6$  PC12 cells (reference ECACC No. 88022) and the unbound sequences are recovered by centrifugation. The latter  
20 sequences are then incubated with  $5 \times 10^6$  adherent PC12 MEN B2 cells, expressing a Ret receptor mutated in the intracellular domain (Ret<sup>M918T</sup>), and the unbound sequences are recovered for the selection phase. This  
25 step makes it possible to select sequences which specifically recognize the PC12 MEN 2A cells expressing the Ret receptor mutated in the extracellular domain (Ret<sup>G634Y</sup>).

The recovered sequences are incubated with  $5 \times 10^6$  PC12  
30 MEN 2A cells for 30 min at 37°C in the presence of non-specific competitive RNA (total yeast RNA, Sigma) and recovered after several washes by total extraction of the RNA (Trizol, Sigma).

35 After an amplification by RT-PCR using the following pair of primers:

sense primer: 5' TAATACGACTCACTATAGGGAGACAAGAATAAACGCTCAA 3' (SEQ ID NO :16)

antisense primer: 5' TCCTGTTGTGAGCCTCCTGTCGTT 3' (SEQ ID NO :17)

and an *in vitro* transcription with mutant T7 polymerase according to the conditions described in Padilla, R et al. (N.A.R., 1999, mentioned above), using a modified  
5 buffer (40 mM Tris, pH 7.5; 6 mM MgCl<sub>2</sub>; 4 mM NaCl; 2 mM spermidine; 10 mM DTT), the process is repeated. All the incubations with the cells are carried out at 37°C in RPMI 1640 culture medium, in order to be as close to physiological conditions as possible.

10

During the selection process, the selection pressure is increased by increasing the number of washes and the amount of non-specific competitive RNA, and decreasing the incubation time and the number of cells exposed to  
15 the aptamers, as illustrated in Table II below.

Table II: Conditions used for the successive selection cycles

Cycle No.	Number (millions) of cells	Nanomoles of 2'-F-Py RNA	Incubation volume (ml)	Concentration (nM) of 2'-F-Py RNA	Incubation period (min)	Number of washes	Competitor added
S1	5	1	3	333	30	1	0
S2	5	3	3	1000	30	1	0
S3	5	3	3	1000	30	2	0
S4	5	0.5	3	167	30	2	0
S5	5	1.5	3	500	15	3	0
S6	5	4	3	1333	15	4	0
S7	5	1.3	3	433	15	3	0
S8	5	2.3	3	767	15	3	0
S9	5	2.5	3	833	15	3	0
S10	5	10	3	3333	15	3	150µg tRNA
S11	5	5.8	3	1933	15	3	150µg tRNA
S12	5	2	3	667	15	4	150µg tRNA
S13	2	1	3	333	15	5	150µg tRNA
S14	2	1	3	333	15	5	150µg tRNA
S15	2	1	3	333	15	5	150µg tRNA

20

In order to follow the evolution of the pool, the appearance of restriction sites containing 4 bases in the population was analyzed by RFLP and reveals the emergence of sequences selected during the SELEX  
25 process, which correspond to specific restriction sites (Bartel DP et al., Science, 1993, 261, 5127, 1411-8).

After fifteen selection cycles, the sequences are

cloned using the TOPO-TA cloning kit (Invitrogen) and analyzed by sequencing.

- Binding assays

5 The binding of the various aptamers (or of the initial pool, as control) to the PC12 MEN 2A cells is carried out in 24-well plates (experiments carried out in triplicate), with 5'-<sup>32</sup>P-labeled RNA. 10<sup>5</sup> cells per well are incubated with various concentrations of aptamers  
10 in 200 µl of RPMI for 10 min at 37°C in the presence of 100 µg/ml of polyinosine, as non-specific competitor. After several washes, the bound sequences are recovered in 350 µl of 0.6% SDS and the amount of radioactivity recovered is related to the number of cells by  
15 measuring the protein content in each well.

The dissociation constants (Kd) and the number of targets (Tmax) for each aptamer are determined:

\* by Scatchard analysis according to the following  
20 equation:

$$[\text{bound aptamer}]/[\text{aptamer}] = -(1/Kd) \times [\text{bound aptamer}] + [Tmax]/Kd$$

\* or by analysis according to the Lineweaver-Burk method, according to the following equation:

$$1/[\text{bound aptamer}] = Kd/([Tmax] \times [\text{aptamer}]) + 1/[Tmax].$$

25

The binding of the individual sequences to the various cell lines is carried out under the same conditions, but at a single concentration of 50 nM.

30 **EXAMPLE 3: Results**

- **SELEX ex vivo: selection of aptamers**

In order to identify the aptamers which recognize the Ret receptor, a modified SELEX protocol was carried out, as specified in example 1:

35 - the method is carried out on intact cells, i.e. a cell line derived from PC12 cells (PC12 MEN 2A), expressing the human mutant receptor Ret<sup>C634Y</sup>;

- 2'-F-pyrimidine RNAs are used to increase the nuclease-resistance of the aptamers present in the

biological fluids;

- two counterselection steps are carried out at each cycle: (1) on PC12 cells which do not express the human Ret receptor and (2) on PC12 MEN 2B cells which express an allele mutated in the intracellular tyrosine kinase domain of the Ret receptor (Ret<sup>M918T</sup>), so as to optimize the selection of aptamers specific for the Ret receptor.

10 The context used made it possible to select aptamers capable of targeting surface epitopes specific for PC12 MEN 2A cells and also aptamers capable of recognizing the extracellular portion of the Ret receptor.

15 This technique has the advantage of avoiding the use of recombinant Ret receptor.

69 clones were obtained and sequenced, from the pool of sequences obtained at the 15th round and bound to the PC12 MEN 2A cells in a saturable manner with a Kd of approximately 100 nM. The results are illustrated in Table III below.

Table III: Results of the selection (the sequences are given without the fixed regions R1 (SEQ ID NO:1) and R2 (SEQ ID NO:2) which border them and allow RT-PCR amplification)



Sequence code (number of clones)	Sequence (R)	binding at 100 nM on PC12 MEN 2A	Kd on PC12 MEN 2A (nM)	number of targets/cell (x 1000)
D14(23)	GGCCATAGCGCACCACCAAGAGCAAAT CCCTAAGCGCGACTCGAGTGAGC	-		
D12(21)	GGGCUUCAUUAAGCUACCGGCCAAC GCAGAAUAGCCUUAAGCCGAGUU	+	67 ± 7	102 ± 25
D30(7)	AGGCGAGCCCGACCACGTAGTATGCT AGACAACAACGCCCGCGTGGTAC	+	74 ± 25	212 ± 27
D71(5)	GGCCCUUAACGCAAAACGAAGGAUCA UCGAUUGAUCGCCUUAUGGGCU	+	53 ± 7	248 ± 42
D42(4)	GACCCGUAUGAAGGUGGCGCAGGACA CGACCGUCUGCAAUGAGCGAGC	-		
D20(2)	GGGCCAATCGAAGCCGTAATTCCCAA ACTAACGTGCAAACTGCACCCGC	-		
D76	GGCTTACACGGAGAAACAAGAGCGCG CCCAAACCTTGATTGACAGTGGCC	-		
D60	CCGACCTGTACAGCAGTTAGTTACAG TTTGAACAACCGCGCTTCGAGC	-		
D32	CCCCGCTTTTGTACGTGATCGAACGCG TATCAGTAACGTGAGCAGTCGAGC	-		
D33	CAAAGCGTGTATTCTCGTGAGCCGACC ATCGTTGCGAACATCCCCGAACG	-		
D87	CCGCGGTCTGTGGGACCTTCAGGATG AAGCGGCAACCCATGCGGGCC	-		
D24	GCGGTATGTAGGGAATAGCACTTTTTT GCGTATACCTACCCGACGCG	+	32 ± 5	102 ± 58
D4	GCGCGGGAATAGTATGGAAGGATACGT ATACCGTGCAATCCAGGGCAACG	+	35 ± 3	110 ± 47

Two sequences (D14 and D12) dominate the selection and constitute more than 50% of the clones. Four other  
5 sequences are less abundant (25% of the clones) and the others are present only once (seven sequences).

Due to the complexity of the target (live whole cells), no similarity was observed between the various aptamer  
10 sequences, except with regard to clones D24 and D4, which share certain motifs and a common structure (see figure 5A, formula II and figures 11 and 12).

The ability of each aptamer to bind to PC12 MEN 2A  
15 cells was tested.

All the sequences found more than once were thus tested, along with sequences present in a lesser amount (including D4 and D24). Despite its abundance, D14 does  
20 not bind the PC12 MEN 2A cells significantly above the background noise.

Other sequences bind the PC12 MEN 2A cells, with a Kd of between 30 and 70 nM.

Most of the aptamers do not bind the parental PC12 cells, rat bladder carcinoma cells (NBTH1) and human HeLa cells.

5

**- Action of the D4 aptamer on the Ret receptor**

The activation of the normal Ret receptor in normal cells (Cn) occurs via interaction with the GRF- $\alpha$  coreceptor for several trophic factors, the most  
10 widespread of which is GDNF (Glial Derived Growth Factor) (figure 4). In cells expressing the normal form of Ret, the cascade is activated only in the presence of GDNF. Inhibition of the phosphorylation of Ret by GDNF in these cells in the presence of D4 is proof that  
15 D4 interacts with the Ret signaling pathway, which confirms the absence of activity of D4 on the activation by another trophic factor, NGF, the activity of which on Erk is not mediated by Ret. The fact that D4 does not have any activity, either, on the  
20 activation of Ret and Erk in MEN 2B cells in which the mutation is intracellular suggests that the interaction between D4 and Ret involves the extracellular portion of Ret, and may be at the dimerization site. However, this is not proved.

25

Very notably, the activity of D4 leads to a reversion of the transformed phenotype on cells in which Ret is constitutively activated. These cells in culture take on a "neuronal-like" morphotype with axonal extensions.  
30 D4, but not D4Sc, which is a destructured D4, having the same chemical composition but without ordering of its sequence in an active structure, induces a very significant reduction in the number of axonal extensions and brings the cell phenotype back to a  
35 nonactivated-cell phenotype, both in a neuroendocrine line (PC 12) and in a fibroblast line (NIH 3T3).

The mutant Ret<sup>C634Y</sup> receptor, expressed by the PC12 MEN 2A cells, forms homodimers at the cell surface, which

leads to constitutive activation of its tyrosine kinase activity (Santoro M et al., Science, 1995, 20, 267, 5196, 381-383) and induces several downstream signaling cascades, including the activation of Erk kinase (Colucci-D'Amato et al., J. Biol. Chem., 2000, **275**, 19306-19314; Jhiang SM, Oncogene, 2000, 19, 5590-5597). Using PC12 cells expressing the Ret<sup>C634Y</sup> oncogene as in vitro cell system, the ability of each aptamer to inhibit the autophosphorylation of the Ret<sup>C634Y</sup> receptor and the downstream signaling dependent on the receptor (figure 6) is evaluated.

The PC12 MEN 2A cells are incubated overnight with one of the following aptamers: D4 (SEQ ID NO:3), D12 (SEQ ID NO:4), D30 (SEQ ID NO:8) and D71 (SEQ ID NO:14), at a final concentration of 150 nM. The cell lysates are analyzed by immunoblotting with anti(Tyr-phosphorylated) Ret antibodies (figure 6B) or anti-phospho-Erk antibodies (figure 6B). As has already been demonstrated (Colucci et al., mentioned above), the levels of phosphorylated Ret receptor and of phosphorylated Erk protein are constitutively high in nontreated PC12 MEN 2A cells, due to the presence of the active Ret<sup>C634Y</sup> allele. Some of the aptamers tested inhibit the autophosphorylation of the Ret<sup>C634Y</sup> receptor and the phosphorylation of the Erk protein which results therefrom, in comparison with the starting combinatorial library and with other aptamers (figure 6A). Using these experimental conditions, the D4 aptamer is found to be the most effective inhibitor and was therefore used for the following studies.

Under these conditions, a dose-response experiment (figure 6B, left) reveals that a concentration of 200 nM of D4 aptamer is sufficient to inhibit the autophosphorylation of the Ret<sup>C634Y</sup> receptor by up to 70% and to very substantially reduce the phosphorylation of the Erk protein. Treatment of the cells for one hour with 200 nM of D4 aptamer is sufficient to

significantly inhibit the autophosphorylation of the Ret<sup>C634Y</sup> receptor and to completely abolish the phosphorylation of the Erk protein (figure 6B, right).

5 In all the experiments, the inhibition of the phosphorylation of the Erk protein is more rapid and more quantitative than the inhibition of the phosphorylation of the Ret<sup>C634Y</sup> receptor, doubtless due to different sensitivities of the two processes for  
10 modifying the tyrosine kinase activity of the Ret receptor.

The predicted secondary structure of the D4 aptamer is illustrated in figure 5A, as is that of the D24  
15 aptamer. Comparison of the two structures suggests that the cell binding is not dependent on the sequence of the tail or of the apical loop. In fact, if the apical loop is replaced with an extra-stable loop comprising four nucleotides (UUGC) or by deleting the nucleotides  
20 represented by R<sub>4</sub> and R<sub>5</sub>, as defined above, no significant difference is observed in binding to the PC12 MEN 2A cells. However, it is the complete D4 aptamer which is the product most active in inhibiting the signaling pathway induced by the Ret<sup>C634Y</sup> receptor. A  
25 2'-F-Py RNA of identical composition but with a destructured sequence (D4Sc) is ineffective both in terms of binding and in terms of inhibition.

The D4 aptamer recognizes the PC12 MEN 2A cells with an  
30 estimated K<sub>d</sub> of 35 nM (figure 5B); furthermore, it recognizes neither the parental PC12 cells, nor rat NBTII cells, nor human HeLa cells, which do not express the Ret receptor.

35 Insofar as the D4 aptamer was selected on cells expressing the mutant Ret<sup>C634Y</sup> receptor, an attempt was made to determine whether the D4 aptamer could also inhibit the wild-type Ret receptor. With this aim, a cell line derived from PC12 cells expressing the wild-

type Ret receptor (PC12/wt) was used. The cells were stimulated with a mixture containing GDNF and soluble GFR $\alpha$ 1 and were treated with the D4 aptamer or with the starting combinatorial library as negative control. As  
5 illustrated in figure 7A, only the D4 aptamer (and not the control combinatorial RNA library) inhibits, substantially, the phosphorylation of the Ret receptor, induced by GDNF (figure 7A), and of the Erk protein (figure 7B). A similar inhibitory effect was observed  
10 with the PC12- $\alpha$ 1/wt cells, a cell line derived from PC12 cells which stably expresses both the human Ret receptor and GFR $\alpha$ 1. It is important to note that the D4 aptamer is inactive on the signaling pathway induced by NGF on the tyrosine kinase receptor TrkA. After  
15 stimulation of the cells with NGF, the treatment with the D4 aptamer does not modify the amount of phospho-Erk, which indicates that the D4-aptamer-induced inhibition of the phosphorylation of the Erk protein is specific for the GDNF-stimulated intracellular  
20 signaling pathway of the Ret receptor (figure 7).

Although the D4 aptamer binds the PC12 MEN 2B cells, treatment of these cells with 200 nM of D4 for one hour (figure 7B) does not interfere with the signaling  
25 pathway induced by the monomeric Ret<sup>M918T</sup> receptor. This confirms that the inhibition of the phosphorylation of the Erk protein by the D4 aptamer is specific for the form activated by dimerization of the Ret receptor. The kinase and biological activities of the Ret<sup>M918T</sup>  
30 receptor, although constitutive, respond to a stimulation with GDNF in the presence of GFR $\alpha$ 1 (Carlomagno F et al., Endocrinology, 1998, 139, 8, 3613-3619). However, in accordance with the inhibition of the activity of the wild-type Ret receptor (Ret wt)  
35 by the D4 aptamer, treatment of the PC12 MEN 2B cells with the D4 aptamer abolishes the overstimulation of the phosphorylation of the Ret receptor and of the Erk protein, dependent on GDNF. These experiments show that the D4 aptamer inhibits the activities of both the Ret

receptor and the Erk protein by direct action on the Ret receptor and not by action on other cell targets, for example tyrosine phosphatase.

5       **- Biological effects of the D4 aptamer on Ret  
receptor-dependent cell differentiation and  
transformation**

The observation that the inhibition occurs in all cases where dimerization of the wild-type (wt) or mutant  
10 receptor is necessary for the signaling pathway also indicates that the dimerization is the target of the action of the D4 aptamer.

With this aim, the axonal extension (or neural crest)  
15 was measured as the reflection of the differentiation in PC12- $\alpha$ 1/wt cells after stimulation with GDNF (see example 1).

The cells are treated with 50 ng/ml of GDNF and the  
20 percentage of cells containing axonal extensions is determined 24 and 48 hours after the treatment as specified in example 1. As illustrated in figure 8, the cells exhibit axonal extension processes in response to exposure to GDNF for two days (figure 8B), compared  
25 with the nonstimulated control cells (figure 8A). The treatment of the cells with the D4 aptamer (figure 8C), but not with the destructured D4 aptamer (D4Sc) used as control (figure 8D), significantly inhibits the number and the length of the axons, perhaps by preventing the  
30 formation of a functional complex between the Ret receptor and GDNF/GFR $\alpha$ 1. In order to biochemically evaluate the differentiation, the VGF levels in cell extracts were determined after 48 hours of treatment. *Vgf* is an early gene which is rapidly induced by NGF  
35 and GDNF in PC12 cells (Salton SR, Mt Sinai J Med., 2003, 70, 2, 93-100). It is observed (figure 8F) that, in the cells treated with GDNF, the expression of VGF is stimulated and in accordance with the phenotypic effects reported above, the treatment with the D4

aptamer, but not the treatment with the destructured D4 aptamer, maintains VGF levels close to basal levels.

After expression of the Ret<sup>C634Y</sup> receptor or of the Ret<sup>M918T</sup> receptor, NIH 3T3 cells are transformed and exhibit considerable changes in their morphology (Santoro et al., Science, 1995, mentioned above). NIH/MEN 2A cells and NIH/MEN 2B cells, which stably express the mutant Ret receptors, are treated with the D4 aptamer for 72 hours and the morphological modifications induced by this aptamer are analyzed. As illustrated in figure 9, the NIH/MEN 2A cells and the NIH/MEN 2B cells have a spindle shape, long protrusions and a highly refringent appearance (figures 9B and 9E, respectively). The NIH/MEN 2A cells treated with the D4 aptamer return to a polygonal and flat morphology similar to that of the parental NIH 3T3 cells (figure 9C), whereas no morphological change is observed in the NIH/MEN 2B cells (figure 9F) or in the NIH-Ras cells. This is in agreement with the previous results which show that the signaling pathway induced by the Ret<sup>C634Y</sup> receptor, but not that induced by the Ret<sup>M918T</sup> receptor is inhibited by the D4 aptamer.

Moreover, the treatment with the destructured D4 aptamer has no effect on the various cell lines (figure 9D). Consequently, treatment of the NIH/MEN 2A cells for 72 hours with the D4 aptamer inhibits the extent of the effects of the activation of the Erk protein by phosphorylation both of the Ret<sup>C634Y</sup> receptor and of the Erk protein.

**EXAMPLE 4: Use of the D4 aptamer for screening products which interact with the Ret receptor or targets which form a complex with the Ret protein on PC12 MEN 2A cells**

In order to validate the use of the D4 aptamer for screening molecules which interact with the Ret

receptor or targets which form a complex with said protein on PC12 MEN 2A cells, two approaches were used:

1 - The D4 aptamer radiolabeled with  $^{32}\text{P}$  was  
5 incubated at 50 nM with monolayers of PC12 MEN 2A cells  
in the presence of 400 nM of various aptamers selected  
either on cells or on the isolated recombinant Ret  
protein. After several washes, the amount of D4 aptamer  
bound is quantified. By comparing the amount of binding  
10 of the D4 aptamer as a function of the molecules used,  
it can be noted that only the D12 aptamer, and  
especially the E38 aptamer, the target of which is the  
Ret receptor, were capable of decreasing the binding of  
the D4 aptamer to the cells. It can therefore be  
15 concluded that these two aptamers bind to a target  
related to the Ret protein, either directly on said  
protein, possibly at the same site as the D4 aptamer,  
or on another target present in a complex with Ret.

20 2 - The E38 aptamer radiolabeled with  $^{32}\text{P}$  was  
incubated at 100 nM with monolayers of PC12 MEN 2A  
cells in the presence of an increasing concentration of  
D4 aptamer. After several washes, the amount of E38  
aptamer bound to the cells was quantified. An  
25 increasing inhibition of the binding of the E38 aptamer  
as a function of the concentration of D4 aptamer can be  
noted. It can therefore be concluded that the E38  
aptamer binds to a target related to the Ret protein,  
either directly on said protein, possibly at the same  
30 site as the D4 aptamer, or on another target present in  
a complex with Ret.